



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the application of: Markus Pompejus, *et al.*

Serial No.: 09/604,693

Filed: June 27, 2000

For: *Corynebacterium Glutamicum Genes Encoding
Proteins Involved in DNA Replication, Protein
Synthesis, and Pathogenesis*

Attorney Docket No.: BGI-130CP

Group Art Unit: 1652

Examiner: Hutson, R.

Commissioner for Patents
Box Sequence Listing
Washington, D.C. 20231

**TRANSMITTAL LETTER FOR COMPACT DISK OF
SUBSTITUTE SEQUENCE LISTING**

Dear Sir:

In response to the Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures mailed from the Patent Office on September 28, 2001, Applicants enclose a Compact Disk (CD) which contains a self-extracting file containing a computer readable form of a Substitute Sequence Listing for the above-referenced patent application filed June 27, 2000. The compact disc is formatted from an IBM-PC and is compatible with MS-Windows. The compact disc contains the following file: seqlistcorr.txt, containing 1,642 kb. This file was created on November 28, 2001. The Substitute Sequence Listing contained on the enclosed compact CD is understood to comply with the requirements of 37 C.F.R. § 1.52 and 37 C.F.R. §1.821-1.825. The material on this diskette is identical in substance to the hard (paper) copy of the Substitute Sequence Listing appearing on substitute pages 1-633 which are submitted herewith, as required by 37 CFR §1.821(f). *No new matter has been added.*

Certificate of First Class Mailing (37 CFR 1.8(a))

I hereby certify that this correspondence is deposited with the United States Postal Service as first class mail in an envelope addressed to: Commissioner for Patents, Washington, D.C. 20231 on:

November 28, 2001

Date

Giulio A. DeConti, Jr., Esq. Reg. No. 31,503

Respectfully submitted,

LAHIVE & COCKFIELD, LLP

Giulio A. DeConti, Jr., Esq.

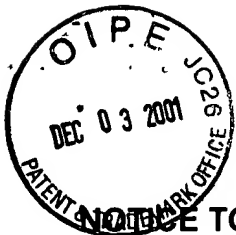
Reg. No. 31,503 for

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Reg. No. 33,505

Attorney for Applicants

Dated: November 28, 2001



Application No.: 09/604,693

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING
NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES**

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

- ☒ 1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
- ☐ 2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
- ☐ 3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
- ☒ 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
- ☐ 5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
- ☐ 6. The paper copy of the "Sequence Listing" is not the same as the computer readable form of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
- ☐ 7. Other: 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the computer readable form is missing.

Applicant Must Provide:

- ☒ An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
- ☐ An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
- ☒ A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

For Rules Interpretation, call (703) 308-4216

For CRF Submission Help, call (703) 308-4212

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